

Figure S3

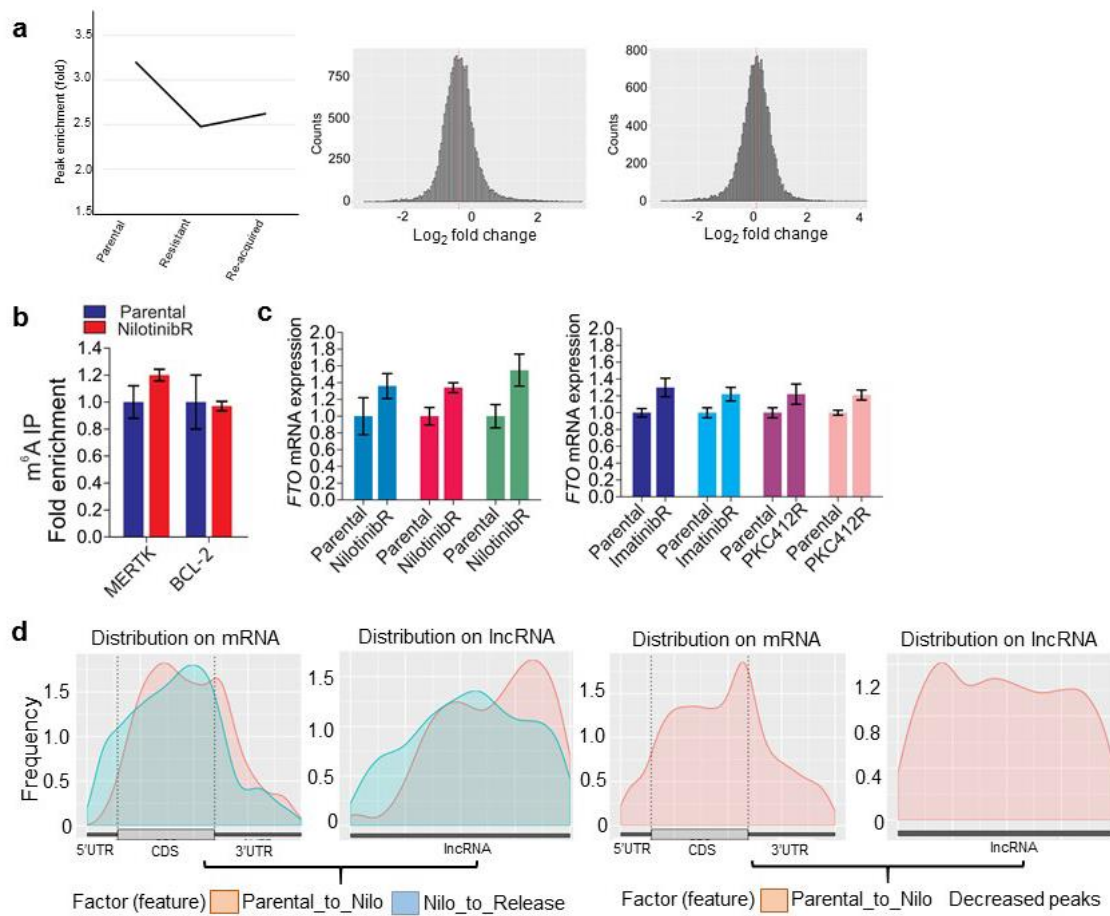


Figure S3. Characterization of *FTO* expression and m⁶A peaks. **a** Analysis of m⁶A peak read accounts. Left, enrichment fold difference was calculated by the total read counts across all peaks in input and IP, and then normalized by total mappable reads for each library; middle and right, plot distribution of log₂ change of enrichment across m⁶A peaks from parental to resistant (middle) or from resistant to reacquired sensitivity to nilotinib (right). The average log₂-transformed normalized signal for the duplicated m⁶A-seq was used to generate a histogram of read counts values. **b** The eluted mRNA from anti-m⁶A immunoprecipitates in parental and resistant cells was subjected to cDNA synthesis followed by qPCR for gene expression using primers outside m⁶A motifs. **c** qPCR of parental and resistant cells for *FTO* expression. **d** Metagene plots of differential

m⁶A peaks across groups. Left, differential peaks (both increased and decreased) of Parental-to-Nilo comparison and Nilo-to-Release comparison; right, peaks showing the decreased enrichment from Parental-to-Nilo comparison.

Nilo, NilotinibR; Release, reacquired sensitivity.